CLEAN VERSION OF THE CLAIMS

- 1. A method for obtaining a biologically active botulinum toxin, comprising the steps of:
- (a) providing a fermentation medium of which not more than about 1 weight percent comprises an animal product selected from the group consisting of an immunoglobulin, a meat product, a meat digest, a meat by product, milk, a dairy product, a dairy digest, blood pooled product, a blood product and an animal protein;
- (b) culturing a Clostridium botulinum bacterium in the fermentation medium under conditions which permit production of a botulinum toxin, wherein the culturing is performed until cell density of the fermentation medium decreases due to cell lysis and;
- (c) recovering a biologically active botulinum toxin from the fermentation medium, wherein the fermentation medium comprises a protein obtained from yeast or from a vegetable, wherein the vegetable is selected from the group consisting of a soy, malt and corn.
- 5. The method of claim 1, wherein in the step of culturing, the culturing is performed until at least 48 hours after initial drop in cell density due to cell lysis.
- 13. A method for making a substantially animal product free pharmaceutical composition in which the active ingredient is a botulinum toxin, the method comprising the steps of:
- (a) obtaining a biologically active botulinum toxin by;
- (i) providing a fermentation medium of which not more than about 1 weight percent is an animal product selected from the group consisting of an immunoglobulin, a meat product, a meat digest, a meat by product, milk, a dairy product, a dairy digest, blood pooled product, a blood product and an animal protein;

- (ii) culturing a Clostridium botulinum in the fermentation medium under conditions which permit production of a botulinum toxin, and;
- (iii) recovering a biologically active botulinum toxin from the fermentation medium:
- (b) formulating the botulinum toxin with a suitable excipient, thereby making a substantially animal product free pharmaceutical composition in which the active ingredient is a botulinum toxin,

wherein the fermentation medium comprises a protein product obtained from yeast or from a vegetable, wherein the vegetable is selected from the group consisting of a soy, malt and corn.

- 14. The method of claim 1, wherein the botulinum toxin is selected from the group consisting of botulinum toxins types A, B, C, D, E, F and G.
- 15. The method of claim 1, wherein the botulinum toxin is a botulinum toxin types A.
- 16. The method of claim 1, wherein the botulinum toxin is a purified botulinum toxin.
- 17. A method for obtaining a biologically active botulinum toxin type A, the method comprising the steps of:
- (a) providing a fermentation medium of which not more than about 1 weight percent comprises an animal product selected from the group consisting of an immunoglobulin, a meat product, a meat digest, a meat by product, milk, a dairy product, a dairy digest, blood pooled product, a blood product and an animal protein;
- (b) culturing a Clostridium botulinum bacterium in the fermentation medium under conditions which permit production of a botulinum toxin, , wherein the culturing is performed until cell density of the fermentation medium decreases due to cell lysis. and;
- (c) recovering a biologically active botulinum toxin from the fermentation medium,

wherein the fermentation medium comprises a protein obtained from yeast or from a vegetable, wherein the vegetable is selected from the group consisting of a soy, malt and corn.

- 18. The method of claim 13, wherein the botulinum toxin is selected from the group consisting of botulinum toxins types A, B, C, D, E, F and G.
- 19. The method of claim 13, wherein the botulinum toxin is a botulinum toxin types A.
- 20. The method of claim 13, wherein the botulinum toxin is a purified botulinum toxin.
- 21. A method for making a substantially animal product free pharmaceutical composition in which the active ingredient is a botulinum toxin type A, the method comprising the steps of:
- (a) obtaining a biologically active botulinum toxin type A by;
- (i) providing a fermentation medium of which not more than about 1 weight percent is an animal product selected from the group consisting of an immunoglobulin, a meat product, a meat digest, a meat by product, milk, a dairy product, a dairy digest, blood pooled product, a blood product and an animal protein;
- (ii) culturing a Clostridium botulinum in the fermentation medium under conditions which permit production of a botulinum toxin type A, and;
- (iii) recovering a biologically active botulinum toxin type A from the fermentation medium;
- (b) formulating the botulinum toxin type A with a suitable excipient, thereby making a substantially animal product free pharmaceutical composition in which the active ingredient is a botulinum toxin type A,

wherein the fermentation medium comprises a protein product obtained from yeast or from a vegetable, wherein the vegetable is selected from the group consisting of a soy, malt and corn.